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		Application Number	0	9/889,106	
TRANSMITTAL FORM (to be used for all carrespondence after Initial filing)		Filing Date	Jı	July 11, 2001 Haile Tecle, et al. 1626	
		First Named Inventor	н		
		Group Art Unit	1		
		Examiner Name	G	Golam M. Shameem	
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	ENCLOS	URES (check all that a	pply)		
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IN THE UNITED STATES PATENT & TRADEMARK OFFICE

APPLICANT: Haile Tecle, et al.

EXAMINER: Golam M. Shameem

SERIAL NO. : 09/889,106

ART UNIT

: 1626

FILED

: July 11, 2001

PAPER NO. : 5

FOR

: 4' HETEROARYL DIARYLAMINES

Response to Restriction Requirement Under 35 U.S.C. 121 and 372

Assistant Commissioner for Patents Washington, D.C. 20231

Dear Sir:

12/16/2002 RHARMON CCCCCCCC 230455 09889106

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110.00 CfThis is in response to the Office Action dated September 30, 2002 (Paper Number 4) in which Restriction was required under 35 U.S.C. 121 and 372.

Under PCT Rule 13.1 and in accordance with 37 CFR 1.499, Applicants were required to elect a single invention of the following inventions:

Group I, claim(s) 1-29, drawn to a compound of the formula (I) wherein, R1 is heterocyclic radical (such as pyrrole, pyrazole), R2 and R3 is independently H, phenyl, C1-4 alkyl, C₃₋₈ alkynyl, C₃₋₈ cycloalkyl, R₄, R₅, and R₆, is independently H, CL, F, or Br, R_A and J are heterocyclic radical (such as pyrrole, pyrazole), R_C, R_D, R_E, R_F and R_G is independently selected from pyrrolidinyl and all other variable substitutions are as defined and one method of use (e.g. treating cancer);

Group II, claim(s) 1-29, drawn to a compound of the formula (I) wherein, R_1 is heterocyclic radical (such as morpholinyl, pyrazinyl), R2 and R3 is independently H, phenyl, C1- $_4$ alkyl, C_{3-8} alkynyl, C_{3-8} cycloalkyl, R_4 , R_5 and R_6 is independently H, CL, F, or Br, R_A and J

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are heterocyclic radical (such as morpholinyl, pyrazinyl), R_C, R_D, R_E, R_F and R_G is independently selected from pyrazinyl and all other variable substitutions are as defined and one method of use (e.g. treating septic shock);

Group III, claim(s) 1-29, drawn to a compound of the formula (I) wherein, R₁ is non-heterocyclic radical (such as H, C₁₋₈ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, phenyl, C₃₋₈ cycloalkyl etc.), R₂ and R₃ is independently H, phenyl, C₁₋₄ alkyl, C₃₋₈ alkynyl, C₃₋₈ cycloalkyl, R₄, R₅ and R₆ is independently H, CL, F, or Br, R_A and J are non-heterocyclic radical (such as H, C₁₋₆ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, phenyl, C₃₋₈ cycloalkyl etc.), R_C, R_D, R_E, R_F and R_G is independently selected from non-heterocyclic radical (such as H, C₁₋₆ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, phenyl, C₃₋₈ cycloalkyl etc.) and all other variable substitutions are as defined and one method of use (e.g. treating cystic fibrosis);

Group IV, claims 30-48, drawn to a method for treating a disease with the compound of the formula (I) wherein, R₁ is heterocyclic radical (such as pyrrole, pyrazole), R₂ and R₃ is independently H, phenyl, C₁₋₄ alkyl, C₃₋₈ alkynyl, C₃₋₈ cycloalkyl, R₄, R₅ and R₆ is independently H, CL, F, or Br, R_A and J are heterocyclic radical (such as pyrrole, pyrazole), R_C, R_D, R_E, R_F and R_G is independently selected from pyrrolidinyl and all other variable substitutions are as defined heterocyclic radical and all other variable substitutions are as defined;

Group V, claims 30-48, drawn to a method for treating a disease with the compound of the formula (I) wherein, R₁ is heterocyclic radical (such as morpholinyl, pyrazinyl), R₂ and R₃ is independently H, phenyl, C₁₋₄ alkyl, C₃₋₈ alkynyl, C₃₋₈ cycloalkyl, R₄, R₅ and R₆ is independently H, CL, F, or Br, R_A and J are heterocyclic radical (such as morpholinyl, pyrazinyl), R_C, R_D, R_E, R_F and R_G is independently selected from pyrazinyl and all other variable substitutions are as defined;

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Group VI, claims 30-48, drawn to a compound of the formula (I) wherein, R₁ is non-heterocyclic radical (such as H, C₁₋₈ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, phenyl, C₃₋₈ cycloalkyl etc.), R₂ and R₃ is independently H, phenyl, C₁₋₄ alkyl, C₃₋₈ alkynyl, C₃₋₈ cycloalkyl, R₄, R₅ and R₆ is independently H, CL, F, or Br, R_A and J are non-heterocyclic radical (such as H, C₁₋₆ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, phenyl, C₃₋₈ cycloalkyl etc.), R_C, R_D, R_E, R_F and R_G is independently selected from non-heterocyclic radical (such as H, C₁₋₆ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, phenyl, C₃₋₈ cycloalkyl etc.) and all other variable substitutions are as defined;

Group VII, claims 49-50, drawn to a method for treating a disease with chemotherapy using the compound of the formula (I) wherein, R₁ is heterocyclic radical (such as pyrrole, pyrazole), R₂ and R₃ is independently H, phenyl, C₁₋₄ alkyl, C₃₋₈ alkynyl, C₃₋₈ cycloalkyl, R₄, R₅ and R₆ is independently H, CL, F, or Br, R_A and J are heterocyclic radical (such as pyrrole, pyrazole), R_C, R_D, R_E, R_F and R_G is independently selected from pyrrolidinyl and all other variable substitutions are as defined;

Group VIII, claims 49-50, drawn to a method for treating a disease with chemotherapy using the compound of the formula (I) wherein, R₁ is heterocyclic radical (such as morpholinyl, pyrazinyl), R₂ and R₃ is independently H, phenyl, C₁₋₄ alkyl, C₃₋₈ alkynyl, C₃₋₈ cycloalkyl, R₄, R₅ and R₆ is independently H, CL, F, or Br, R_A and J are heterocyclic radical (such as morpholinyl, pyrazinyl), R_C, R_D, R_E, R_F and R_G is independently selected from pyrazinyl and all other variable substitutions are as defined; or

Group IX, claims 49-50, drawn to a method for treating a disease with chemotherapy using the compound of the formula (I) wherein, R_I is non-heterocyclic radical (such as H, C₁₋₈ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, phenyl, C₃₋₈ cycloalkyl etc.), R₂ and R₃ is independently H, phenyl, C₁₋₄ alkyl, C₃₋₈ alkynyl, C₃₋₈ cycloalkyl, R₄, R₅ and R₆ is independently H, CL, F, or Br, R_A and J are non-heterocyclic radical (such as H, C₁₋₆ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, phenyl, C₃₋₈ cycloalkyl etc.), R_C, R_D, R_E, R_F and R_O is independently selected from non-heterocyclic

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radical (such as H, C₁₋₆ alkyl, C₃₋₈ alkenyl, C₃₋₈ alkynyl, phenyl, C₃₋₈ cycloalkyl etc.) and all other variable substitutions are as defined.

The Applicants hereby elect Group III for prosecution on the merits, without traverse.

The Applicant hereby select the treatment of cancer as the one method of use to accompany the compounds of Group III. Finally, the Applicants respectfully reserve the right to file a Divisional application(s) to the non-elected claims.

The Applicants submit that this application is now in condition for allowance, which allowance is respectfully solicited.

Respectfully submitted,

Dated: December 2, 2002

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